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(54) Title: ANTI-INFLAMMATORY BIOACTIVE GLASS PARTICULATES (57) Abstract Compositions and methods for systemically minimizing the inflammatory effects of TNF- α are disclosed. The compositions include particles of bioactive glass with a particle size less than about 20 μm , alone or in combination with anti-inflammatory agents and other therapeutic agents. The compositions can include an appropriate carrier for oral, intramuscular, intraperitoneal or intravenous administration. When administered to a patient, the particles bring about elevated levels of IL-6 and do not cause elevated levels of TNF- α .		

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ANTI-INFLAMMATORY BIOACTIVE GLASS PARTICULATES

FIELD OF THE INVENTION

The present invention relates to compositions and methods for use in
5 the transient suppression of the inflammatory response by suppressing
plasma levels of tissue necrosis factor-alpha and increasing plasma levels of
interleukin-6.

BACKGROUND OF THE INVENTION

10 When an injury occurs, cell damage initially comes from the
precipitating event, such as a cut, resulting in ruptured cells and severed or
crushed capillaries and other blood vessels. The interruption of blood flow
results in anoxia, causing the death of additional cells. The wound site
quickly fills with dead and dying cells, extracellular substances (collagen,
15 elastic fibers, fat and ground substances), extravasated blood, and possibly
bacteria and viruses introduced during the injury.

Tissue damage is not restricted to the initial area of injury, and may
increase over the next several hours or days as a result of the release of
20 lysosomal enzymes from the injured cells or as a consequence of inflammation
(swelling) and/or infection. (See Reese et al., Role of Fibronectin in Wound
Healing, the subject matter of which is hereby incorporated by reference.)
The inflammatory response is one of the normal stages of wound healing,
and is necessary for subsequent phases of healing.

25

Inflammation is a vital process necessary for an organism to survive
an external insult, such as a wound or burn. However, if unchecked,
inflammation can have harmful consequences. For example, many chronic
and even life-threatening disorders, such as asthma, rheumatoid arthritis,

lung fibrosis, peritoneal adhesions, hypersensitivity and autoimmune diseases are a result of an uncontrolled inflammatory response. An unresolved inflammation in the lung resulting from bacterial infection (i.e., pneumonia) may eventually lead to extensive tissue damage and a chronic lung abscess. Inflammation of the peritoneal cavity, for example, and the resulting adhesions following abdominal surgery is a major cause of infertility in women. Asthma is an often life-threatening disorder which results from an inadvertently stimulated inflammatory response in the lungs. An excessive inflammatory response can cause extensive swelling, which can lead to additional injury as a result of anoxia. Pain results from a combination of kinins and the effect of lysozymes and pressure from the swelling on nerve endings. Unchecked, the inflammatory response can set off a neural feedback loop and cause hyperalgesia, a phenomenon in which the surrounding area of injury remains painful. Accordingly, there is a great interest in the medical community to develop anti-inflammatory agents.

Many anti-known inflammatory compositions reduce the inflammatory response, but are also immunosuppressive. For example, corticosteroids are potent anti-inflammatory agents, but are associated with T-cell suppression and increased infections. Interleukin-10 (as well as IL-4 and IL-3 to lesser extents) are broadly acting anti-inflammatory agents, but are associated with decreased cell mediated immune functions.

It would be advantageous to provide compositions and methods which provide protection from adverse effects associated with inflammation, preferably without unnecessary immunosuppression. The present invention provides such compositions and methods.

SUMMARY OF THE INVENTION

Compositions and methods for providing protection from adverse effects associated with inflammation are disclosed. The compositions and methods can suppress plasma concentrations of tissue necrosis factor-alpha (TNF- α) while increasing plasma concentrations of interleukin-6 (IL-6).

The compositions include non-interlinked particles of bioactive glass with a size less than about 20 μm , alone or in combination with an additional anti-inflammatory agent, and optionally include other therapeutic agents. Formulations including the composition and a suitable carrier, preferably for oral, intramuscular, intraperitoneal or intravenous administration, are also disclosed.

The composition can be administered orally, intramuscularly, intraperitoneally or intravenously to provide systemic relief from the adverse effects associated with inflammation, for example, the effects of excess TNF- α .

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a graph showing the number of cells (cells/ml $\times 10^4$) for mice injected intraperitoneally with control and with bioactive glass, as described in Example 1.

Figure 2 is a graph showing the concentration of TNF- α (pg/ml) for mice injected intraperitoneally with control and with bioactive glass, as described in Example 1.

Figure 3 is a graph showing the concentration of IL-6 (pg/ml) for mice injected intraperitoneally with control and with bioactive glass, as

described in Example 1.

Figure 4 is a graph showing the concentration of IL-6 (pg/ml) for mice injected intraperitoneally with varying doses of bioactive glass, as described in Example 2.

Figure 5 is a graph showing the concentration of TNF- α (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS/D-galactosamine, as described in Example 2.

Figure 6 is a graph showing the concentration of TNF- α (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS, as described in Example 2.

Figure 7 is a graph showing the concentration of IL-1 α (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS, as described in Example 2.

Figure 8 is a graph showing the concentration of IL-6 (pg/ml) for mice injected intraperitoneally with bioactive glass and LPS, as described in Example 2.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Compositions and methods for providing protection from adverse effects associated with inflammation are disclosed. The compositions and methods can suppress plasma concentrations of tissue necrosis factor-alpha (TNF- α) while increasing plasma concentrations of interleukin-6 (IL-6).

The compositions can be administered orally, intramuscularly,

intraperitoneally or intravenously to provide systemic relief from the adverse effects associated with inflammation, for example, the effects of excess TNF- α . The bioactive glass is bioactive *in vivo*, and is not pro-inflammatory. It does not cause the recruitment of PMNs and does not
5 stimulate TNF- α secretion. It stimulates IL-1 β release very modestly, and induces an IL-6 response.

IL-6 is a unique cytokine with pro- and anti-inflammatory properties. It is the primary hepatic acute phase response inducer (a pro-inflammatory
10 effect). It suppresses TNF- α production by macrophages (an anti-inflammatory effect), promotes B-cell proliferation and promotes a Th₂ response, and is mitogenic for some cell types. It has been shown to reduce collagen induced arthritis (Immunology, 95(1):31 (1998)) and to reduce monocyte-mediated TNF production in response to LPS (Exp. Cell Res.,
15 215(1):51-56 (1994) and Am. J. Physiol. 267 (4pt.1):L442-446 (1994)).

The compositions include non-interlinked particles of bioactive glass with a size less than about 20 μ m, alone or in combination with an additional anti-inflammatory agents, and optionally include other therapeutic agents.
20 Formulations including the composition and a suitable carrier, preferably for oral or intravenous administration, are also disclosed.

The normal inflammatory effect is generally regarded as being ligand/receptor controlled. Not being bound to any particular theory or
25 mechanism, it is believed that the rate and amount of ions generated by the absorption of the small particles of bioactive glass has an effect on the pro-inflammatory receptors, which is responsible for the suppression of the cytokines involved in the inflammatory process. Further, the surface of the small particles is micro-porous, and may adsorb the cytokines, rendering

them inactive in the inflammatory process.

The compositions and methods described herein are advantageous
5 because they reduce the inflammatory response, but are not broadly
immunosuppressive. There does not appear to be a typical dose dependency
in terms of particles of bioactive glass and IL-6 response. The response is
independent of dose up to extremely high levels of particulate, for example,
over XXX mg injection. Further, the bioactive glass particles appear to elicit
10 a specific IL-6 response, transient in nature, lasting on the order of several
hours. Pre-exposure to bioactive glass particles results in a significantly
attenuated pro-inflammatory IL-6 cytokine response that results in the
suppression of a typical inflammatory response caused by injection of an
endotoxin after the injection of the bioactive glass particles.

15

The terms “wound” and “burn,” collectively referred to herein as
“injury” have their usual meanings. “Wound” is intended to include wounds
caused by surgical procedures. “Normal” is used in the sense it is usually
used in the medical arts. “Medical practitioner” means one of ordinary skill
20 in the art wound and burn treatment. Typically, this person is a physician,
nurse, dentist, or paramedic.

I. Bioactive Glass

Compositions including non-interlinked particles of bioactive glass
25 with an average diameter of less than about 20 μm , alone or in combination
with anti-inflammatory agents, can be used for the methods described herein.

Very small particulate bioactive glass has the property of exerting an
anti-inflammatory effect when administered systemically. It appears that the

bioactive glass suppresses the production of tissue necrosis factor alpha (TNF- α) and interleukin-6 (IL-6). TNF- α is a powerful pro-inflammatory cytokine that not only participates in the normal inflammatory response, but is also implicated in myocardial dysfunction and cardiomyocyte death in
5 ischemia-reperfusion injury, sepsis, chronic heart failure, viral myocarditis and cardiac allograft rejection, as well as a host of other inflammatory disorders. Accordingly, by suppressing the production of TNF- α , the compositions reduce the likelihood of these disorders occurring.

10 The preferred size range for the bioactive glass, for this embodiment, is such that the particles do not physically obstruct vascular, lymph or pulmonary pathways as the particles pass through the body. As the particles are less than about 20 microns in size, they avoid phagocytosis and uptake by the reticuloendothelial system. This is in stark contrast to small particles of
15 other materials, such as talcum, asbestos, silicone and metal debris, which are known to be strongly pro-inflammatory. The particles are of a suitable size for intravenous administration.

As used herein the terms "bioactive glass" or "biologically active
20 glass" mean an inorganic glass material having an oxide of silicon as its major component and which is capable of bonding with growing tissue when reacted with physiological fluids.

Bioactive glasses are well known to those skilled in the art, and are
25 disclosed, for example, in *An Introduction to Bioceramics*, L. Hench and J. Wilson, eds. World Scientific, New Jersey (1993), the contents of which are hereby incorporated by reference.

The glass preferably includes between 40 and 86% by weight of

silicon dioxide oxide (SiO_2), between about 0 and 35% by weight of sodium oxide (Na_2O), between about 4 and 46% by weight calcium oxide (CaO), and between about 1 and 15% by weight phosphorus oxide (P_2O_5). More preferably, the glass includes between 40 and 60% by weight of silicon dioxide oxide (SiO_2), between about 5-30% by weight of sodium oxide (Na_2O), between about 10 and 35% by weight calcium oxide (CaO), and between about 1 and 12% by weight phosphorus oxide (P_2O_5). The oxides can be present as solid solutions or mixed oxides, or as mixtures of oxides.

10 CaF_2 , B_2O_3 , Al_2O_3 , MgO and K_2O may be included in the composition in addition to silicon, sodium, phosphorus and calcium oxides. The preferred range for B_2O_3 is between 0 and 10% by weight. The preferred range for K_2O is between 0 and 8% by weight. The preferred range for MgO is between 0 and 5% by weight.

15 The most preferred glass is Bioglass®TM (a trademark of University of Florida), which has a composition including about 45% by weight silicon dioxide, about 24.5% by weight sodium oxide, about 6% by weight phosphorus oxide, and about 24.5% by weight calcium oxide. Another preferred material is hydroxyapatite.

20 Particulate, non-interlinked bioactive glass is preferred in the present invention. That is, the glass is in the form of small, discrete particles, rather than a fused matrix of particles or a mesh or fabric (woven or non-woven) of glass fibers. Note that under some conditions the discrete particles of the present invention may tend to cling together because of electrostatic or other forces but are still considered to be non-interlinked. The particle size is less than about 20 microns, preferably 10 microns or less, more preferably less than about 5 microns, and ideally, less than about 2 microns.

The glass composition can be prepared in several ways, to provide melt-derived glass, spun fibers of sol-gel derived glass, and sintered glass particles. The sintered particles may be in sol-gel derived, or pre-reacted melt derived form. Sol-gel derived glass is generally prepared by
5 synthesizing an inorganic network by mixing metal alkoxides in solution, followed by hydrolysis, gelation, and low temperature (600-900°C) firing to produce a glass. Melt derived glass is generally prepared by mixing grains of oxides or carbonates, melting and homogenizing the mixtures at high temperatures, typically between about 1250 and 1400°C. The molten glass
10 can be fritted and milled to produce a powder or casted into steel or graphite molds to make bulk implants.

The glass composition is preferably melt-derived. In each preparation, it is preferred to use reagent grade glass, especially since the
15 glass is used to prepare materials which ultimately may be implanted in a human. Other compositions which biodegrade and release the same ions as the bioactive glasses described above can also be used, provided they do not elicit an inflammatory response and that they also elicit enhanced IL-6 production.

20

A. Melt Derived Glass

A melt-derived glass composition can be prepared, for example, by preparing an admixture of the individual metal oxides and other components used to prepare the glass composition, blending the admixture, melting the
25 admixture, and cooling the mixture. The melting temperature is determined in large part by the glass composition, and ranges, for example, from about 900-1500°C, preferably between about 1250 and 1450°C. The melt is preferably mixed, for example, by oxygen bubbling, to ensure a thorough homogenation of the individual components.

The mixture can be cooled, for example, by adding the molten admixture to a suitable liquid, such as deionized water, to produce a glass frit. Porosity can be introduced by grinding the glass into a powder, admixing the powder with a foaming agent, and hot pressing the mixture under vacuum and elevated temperature. The particle size of the glass powder is between about 40 and 70 μm , the vacuum is preferably less than 50 MPa, and the hot pressing is preferably performed at a temperature above 400°C, preferably between about 400 and 500°C. Suitable foaming agents include compounds which evolve carbon dioxide and/or water at elevated temperatures, for example, metal hydroxides, metal carbonates, and peroxides, such as hydrogen peroxide. Preferred metal carbonates are sodium bicarbonate, sodium carbonate and calcium carbonate. The foaming agents are preferably added in a range of between about 1-5, more preferably 2-3 percent by weight of the glass powder. The preparation of melt-derived porous glass is described, for example, in U.S. Patent No. 5,648,301 to Ducheyne and El Ghannam, the contents of which are hereby incorporated by reference.

B. Sintered Glass Particles

Glass can be sintered using known methodology. In one embodiment, an aqueous slurry of the glass powder and a foaming agent with a suitable binder, such as polyvinyl alcohol, is formed. The slurry is then poured into a mold, allowed to dry, and sintered at high temperatures. These temperature may range, depending on the glass composition and foaming agent used, between about 500 and 1000°C, more preferably between about 600 and 800°C.

C. Spun Fibers of Sol-gel Derived Glass

It is known in the art to control the heat treatment cycle of glass gels

to control the pores and interpores of the material to create a porous glass material. Suitable pore diameters are between 20 and 180Å, suitable pore volumes are between 40 and 52 cc/g, and suitable surface areas are between 75 and 350 m²/g. Since a pore diameter larger than 0.1 microns is difficult to achieve using this method, the sintering and foaming processes described
5 herein are generally more preferred.

D. Leaching of the Porous Material

To aid in preparing glass compositions with high porosity, the glass composition can include a material which can be preferably leached out of
10 the glass composition, and, in doing so, provide the composition with high porosity. For example, minute particles of a material capable of being dissolved in a suitable solvent, acid, or base can be mixed with or melted into the glass, and subsequently leached out. The resulting voids have roughly
15 the same size as the particle that was leached out. In the case of a material which is part of a melt-derived glass composition, the size of the pores and degree of porosity depends on the amount of added material relative to the amount of glass. For example, if the leached material constituted about 80% of the glass, then the glass would be approximately 80% porous when the
20 material was leached out. When leaching the glass composition, care should be taken not to leach out those components which add to the bioactivity of the glass, i.e., the calcium and phosphorus oxides.

II. Formulations including Bioactive Glass

25 The bioactive glass particulates are preferably administered in a formulation that includes an acceptable carrier for the mode of administration. Suitable pharmaceutically acceptable carriers are known to those of skill in the art. The formulations can optionally include other therapeutically active ingredients, such as antibiotics, antivirals, healing

promotion agents, anti-inflammatory agents, immunosuppressants, growth factors, anti-metabolites, cell adhesion molecules (CAMs), antibodies, vascularizing agents, anti-coagulants, and anesthetics/analgesics.

- 5 The carrier must be pharmaceutically acceptable in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof. The formulations can include carriers suitable for oral, rectal, topical or parenteral (including subcutaneous, intramuscular and intravenous) administration. Preferred carriers are those suitable for oral or
10 parenteral administration.

- Formulations suitable for parenteral administration conveniently include a sterile aqueous preparation of the bioactive glass. Such formulations may conveniently contain distilled water, 5% dextrose in
15 distilled water or saline. The formulations may be in the form of a powdered formulation including the bioactive glass particulates and salt, dextrose, buffers and the like, which can be dissolved in water immediately prior to administration.

- 20 For enteral administration, the bioactive glass particulates can be incorporated into an inert carrier in discrete units such as capsules, cachets, tablets or lozenges, each containing a predetermined amount of the particles; as a powder or granules; or a suspension or solution in an aqueous liquid or non-aqueous liquid, *e.g.*, a syrup, an elixir, an emulsion or a draught.
25 Suitable carriers may be starches or sugars and include lubricants, flavorings, binders, and other materials of the same nature.

 A tablet may be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by

compressing in a suitable machine the active compound in a free-flowing form, e.g., a powder or granules, optionally mixed with accessory ingredients, e.g., binders, lubricants, inert diluents, surface active or dispersing agents. Molded tablets may be made by molding in a suitable machine, a mixture of the powdered active compound with any suitable carrier.

A syrup or suspension may be made by adding the active compound to a concentrated, aqueous solution of a sugar, e.g., sucrose, to which may also be added any accessory ingredients. Such accessory ingredients may include flavoring, an agent to retard crystallization of the sugar or an agent to increase the solubility of any other ingredient, e.g., as a polyhydric alcohol, for example, glycerol or sorbitol.

In addition to the aforementioned ingredients, the formulations may further include one or more optional accessory ingredient(s) utilized in the art of pharmaceutical formulations, e.g., diluents, buffers, flavoring agents, binders, surface active agents, thickeners, lubricants, suspending agents, preservatives (including antioxidants) and the like.

20

Other Therapeutic Agents

In addition to bioactive glass particles, the formulations can include other therapeutic agents such as antibiotics, antivirals, healing promotion agents, anti-inflammatory agents, immunosuppressants, growth factors, anti-metabolites, cell adhesion molecules (CAMs), bone morphogenic proteins (BMPs), vascularizing agents, anti-coagulants, and topical anesthetics/analgesics.

The antibiotics can be topical antibiotics suitable for skin treatment.

Examples of such antibiotics include but are not limited to: chloramphenicol, chlortetracycline, clyndamycin, clioquinol, erythromycin, framycetin, gramicidin, fusidic acid, gentamicin, mafenide, mupirocin, neomycin, polymyxin B, bacitracin, silver sulfadiazine, tetracycline and
5 chlortetracycline.

Suitable antivirals include topical antivirals, such as acyclovir and gancyclovir. Suitable anti-inflammatory agents include corticosteroids, hydrocortisone and nonsteroidal antinflammatory drugs. Suitable growth
10 factors include basic fibroblast growth factor (bFGF), epithelial growth factor (EGF), transforming growth factors α and β (TGF α and β), platelet-derived growth factor (PDGF), and vascular endothelial growth factor/vascular permeability factor (VEGF/VPF)). Suitable topical anesthetics include benzocaine and lidocaine.

15 An additional anti-inflammatory agent which can be used in combination with the bioactive glass particles is β -glucan. β -glucan is a polysaccharide with anti-inflammatory properties, and, like the bioactive glass particles described herein, increases IL-6 release and decreases TNF- α and IFN γ release.
20

III. Methods of Reducing Inflammation

The compositions can be used to prophylactically or therapeutically to reduce inflammation in a patient. Overly acute or chronic inflammation
25 can result in various disease states in a patient, for example, arthritis and tendonitis, pulmonary disorders such as asthma and emphysema, and post-surgical (peritoneal) adhesions.

The particles of bioactive glass are believed to modify the inflammatory response in the local microenvironment by altering the synthetic properties of resident macrophages and other recruited inflammatory cells. The particles induce macrophage tolerance by eliciting a cytokine response (IL-6 based) that acts in an autocrine and paracrine fashion.

Particles of bioactive glass can be delivered by intravenous, intramuscular, or intraperitoneal injection to provide systemic (and possibly local when administered intramuscularly or intraperitoneally) anti-inflammatory effects. These effects can be therapeutic and/or prophylactic. For example, systemic delivery of bioactive glass can be effective in reducing the onset of inflammation brought on by external challenge. Particles of bioactive glass can also be administered orally.

Systemic administration of the bioactive glass particles can lower production of TNF- α and also increase production of IL-6. The effect of this is to modify the pro-inflammatory response, and will be useful in treating or preventing sepsis, systemic inflammatory response syndromes, adult respiratory distress syndrome, ocular injury, surgical wound healing and adhesion formation, and delayed cutaneous wound healing. Other disorders which are, at least in part, attributable to an exaggerated inflammatory response, such as ARDS, pancreatitis, viral hepatitis, hemorrhagic shock, ischemia/reperfusion injury, peritoneal adhesions, and chronic inflammation, such as delayed wound healing and rheumatoid arthritis, can be treated with systemic administration of bioactive glass particles.

In one embodiment, the particles are administered locally, for example, by inhalation, by spraying (in the form of an aerosol) or by mixing

the particles with a gel (for example, a biocompatible hydrogel), creme or other aqueous or non-aqueous carrier, and applying or injecting the composition subcutaneously at a site at which surgery is to be performed at a later time. The compositions can advantageously include an anesthetic. The presence of the particles at the proposed surgical site can lead to decreased inflammation resulting from the surgery.

The present invention will be more clearly understood with reference to the following non-limiting examples.

10

EXAMPLES

Example 1: Intraperitoneal Administration of Bioactive Glass to Mice

Ten mice were injected intraperitoneally with 25 mgs of bioactive glass (45S5) with a particle size less than about 20 μm in a total volume of 1 ml (0.5 ml fetal calf serum and 0.5 ml phosphate-buffered saline) with a result pH of 9.6. An additional ten mice received the carrier (0.5 ml fetal calf serum and 0.5 ml phosphate-buffered saline) with the pH unadjusted. Two hours later, the mice were euthanized and the peritoneal contents were washed with 3 ml of physiologic saline. Peritoneal white cell count was performed with a hemocytometer, and peritoneal TNF- α and IL-6 determined by ELISA.

The procedures were performed without incident. The animals appeared grossly normal for the two hour post-injection period. At sacrifice, peritoneal white cell count in mice receiving the bioactive glass particles was not different from peritoneal wash fluid from mice receiving only the carrier. The proinflammatory cytokine TNF- α was not detected in any of the samples. Peritoneal IL-6 concentrations, however, were increased 25 fold

from approximately 80 pgs/ml in the carrier-treated mice to over 2,000 pgs/ml in the bioactive glass-treated mice. The cell count, TNF- α concentration and IL-6 concentration from the peritoneal washings for the bioactive glass-treated mice and the control mice are shown in Figures 1-3.

5

In conclusion, the bioactive glass is bioactive when administered intraperitoneally. The bioactive glass is not acting as a classical irritant. A chemical irritation or a direct inflammatory agent like endotoxin or TNF- α would have resulted in a significant and rapid neutrophil influx, which was not observed. The bioactive glass was not directly pro-inflammatory, since no TNF- α response was elicited. A significant IL-6 response was observed, but IL-6 has both pro-inflammatory (hepatic acute phase induction) and anti-inflammatory properties (delays apoptosis, suppresses macrophase TNF and IL-1 production). Since no inflammatory cell influx was observed, the enhanced IL-6 production must have been secondary to resident macrophages and fibroblasts. The results are consistent with bioactive glass-mediated stimulation of resident cell IL-6 synthesis, which represents a new anti-inflammatory property.

20 Example 2: Anti-Inflammatory Effect of Bioactive Glass in a Mouse Endotoxemia Model

Bioactive glass particles with a particle size of 5 μ m were injected to C57B1/6 mice. The mice were injected ip with 3, 6, 12, 25 and 50 mg of bioactive glass particles or buffer. The animals were sacrificed and TNF- α , IL-1 α and IL-6 levels were determined in the peritoneal lavage fluid at 2 hours (Figure 4). In a second study, mice were injected with 12 mg of bioactive glass particles or buffer ip and then challenged ip with 1 μ g of LPS and 8 mg of D-galactosamine (Dgal) 6 hours after the bioactive glass

particles were administered. Blood samples were taken 90 minutes after Dgal/LPS injection for determination of TNF- α . Lastly, a sublethal dose of LPS (100 ng) was given ip 2 hours after an ip injection of 3 mg of 5 μ m particles of bioactive glass. The mice were bled, sacrificed and lavaged 2
5 hours after the LPS administration.

Results: In the first study, all doses of bioactive glass were found to induce a significant IL-6 response (range 4,584-23,112 pg/ml bioactive glass, $1,017 \pm 445$ pg/ml \pm SEM buffer); however, the bioactive glass did not induce TNF- α or IL-1 α production in the peritoneal lavage fluid. In the second study,
10 mice that received bioactive glass particles prior to Dgal/LPS had significantly lower plasma TNF- α than did controls (bioactive glass: $13,047 \pm 4,126$ pg/ml, control: $34,813 \pm 4,902$ pg/ml, $p < 0.0038$) (Figure 5). Finally, peritoneal IL-6 and TNF- α were reduced in response to LPS by pretreatment
15 with bioactive glass (IL-6 $24,452 \pm 6,673$ pg/ml and control: $53,330 \pm 2,586$ pg/ml, bioactive glass: TNF- α - 26 ± 5 pg/ml and control: 91 ± 49 pg/ml) (Figures 6-8).

Conclusion: Bioactive glass particles are a bioactive substance that when
20 administered alone appear to elicit a significant initial IL-6 response without concurrent expression of TNF- α or IL-1 α .

We claim:

1. A composition comprising particles of bioactive glass with a particle size less than about 20 μm in diameter and a suitable carrier for oral, intramuscular, intraperitoneal or intravenous administration.
5
2. The composition of claim 1, wherein the carrier is suitable for intravenous administration.
3. The composition of claim 1, wherein the carrier is suitable for
10 intramuscular or intraperitoneal administration.
4. The composition of claim 1, additional comprising one or more therapeutic agents.
- 15 5. The composition of claim 4, wherein one or more therapeutic agents are selected from the group consisting of healing promotion agents, growth factors, anti-inflammatory agents, and anesthetics.
6. The composition of claim 1 wherein the glass includes between
20 about 40 and 86 percent by weight of SiO_2 , between about 0 and 30 percent by weight of Na_2O , between about 4 and 46 percent by weight of CaO and between about 1 and 15 percent by weight of P_2O_5 .
7. The composition of claim 1, wherein the bioactive glass has a
25 particle size range less than about 2 microns.
8. A method for systemically minimizing the production of $\text{TNF-}\alpha$ caused by an inflammatory response in a patient comprising orally or

intravenously administering an effective TNF- α lowering amount of bioactive glass particles with a size less than about 20 μm to the patient.

9. The method of claim 8 wherein the systemic minimization of TNF- α is used prophylactically or therapeutically to prevent or treat a disease state selected from the group consisting of peritoneal adhesions, septic shock, cardiac allograft rejection, systemic inflammatory response syndromes, adult respiratory distress syndrome, ocular injury, delayed cutaneous wound healing, ARDS, pancreatitis, viral hepatitis, hemorrhagic shock, ischemia, reperfusion injury, and rheumatoid arthritis.

10

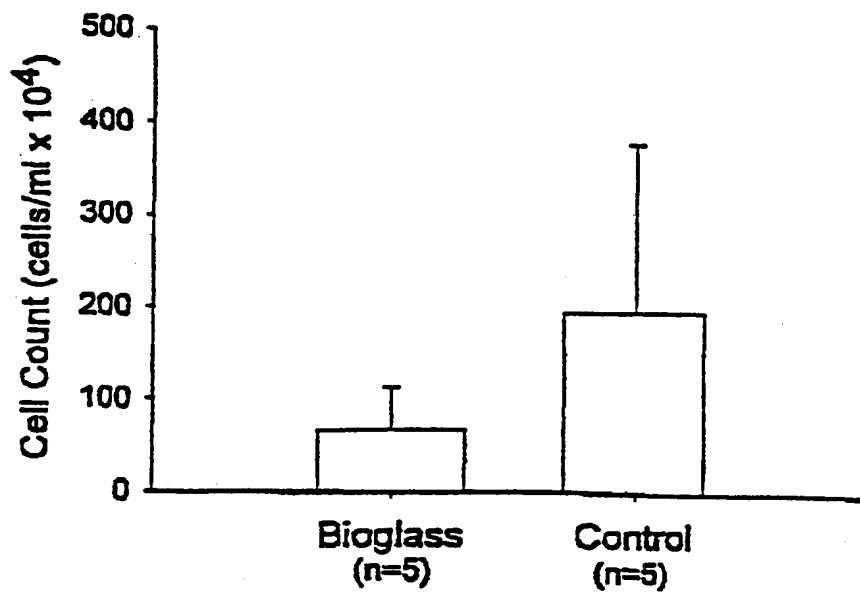
10. A method for systemically increasing IL-6 levels in a patient, comprising administering to the patient an effective, IL-6 increasing amount of bioactive glass particles with a size less than about 20 μm .

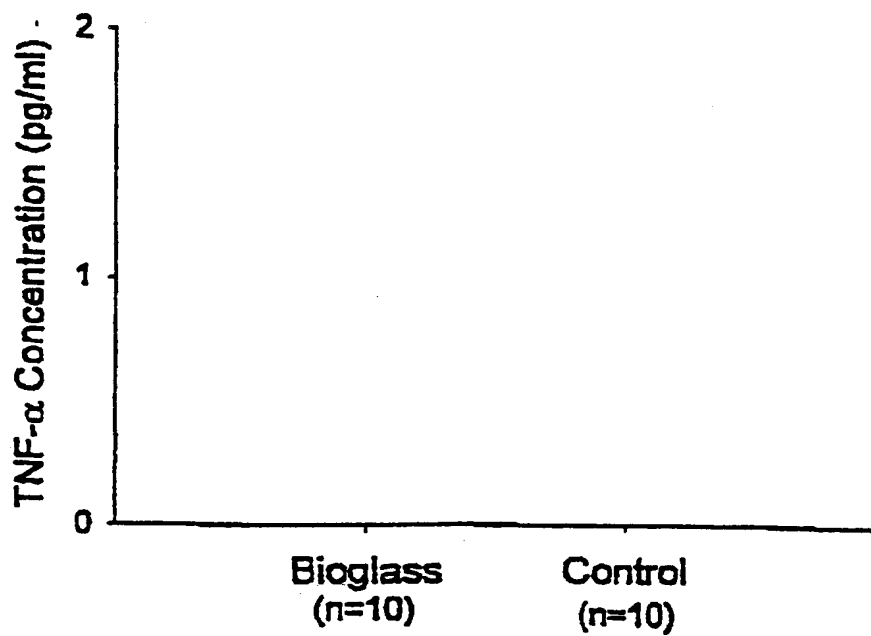
15

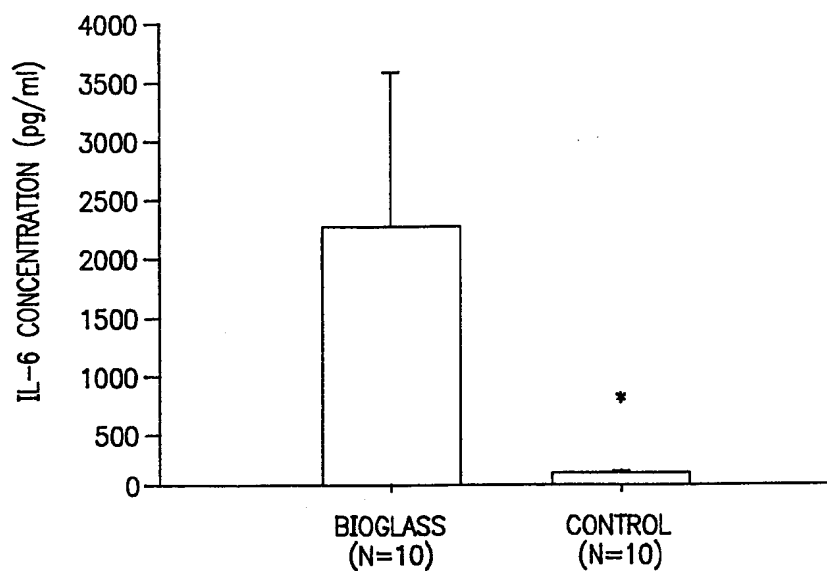
11. A composition comprising particles of a material with a particle size less than about 20 μm which biodegrades, produces elevated serum concentrations of calcium and phosphorous ions, does not cause elevated plasma TNF- α concentrations, and does cause elevated plasma IL-6 concentrations, in combination with a suitable carrier for oral, intramuscular, intraperitoneal or intravenous administration.

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**Fig. 1**

**Fig. 2**



*($p < 0.0001$ BY MANN-WHITNEY RANK SUM TEST)

FIG. 3

IL-6 LEVELS IN PERITONEAL LAVAGE FLUID
FOLLOWING VARIOUS DOSES OF ip BIOGLASS

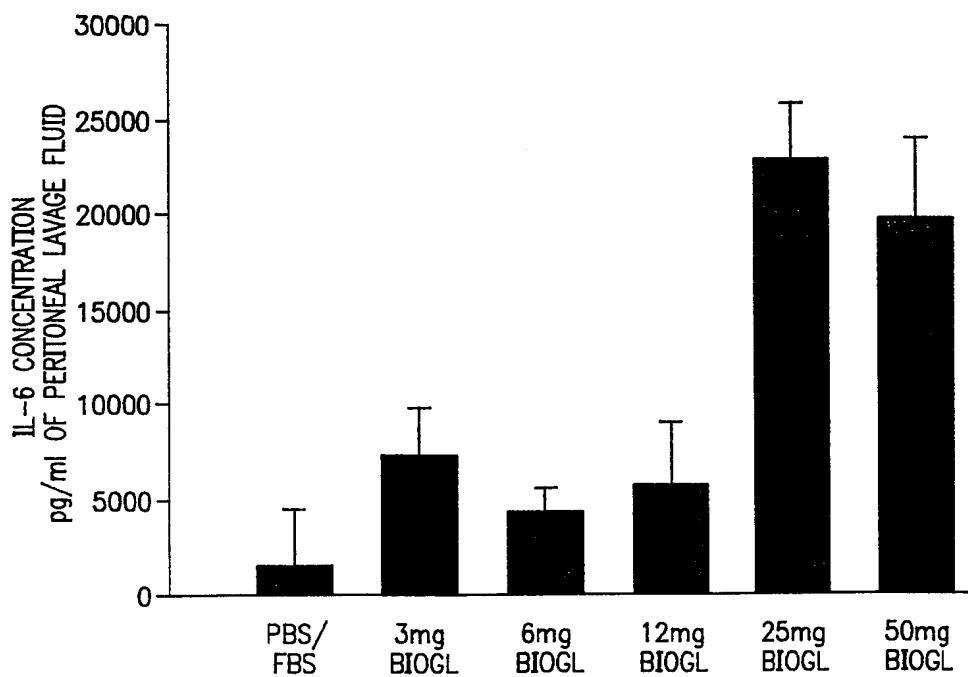


FIG. 4

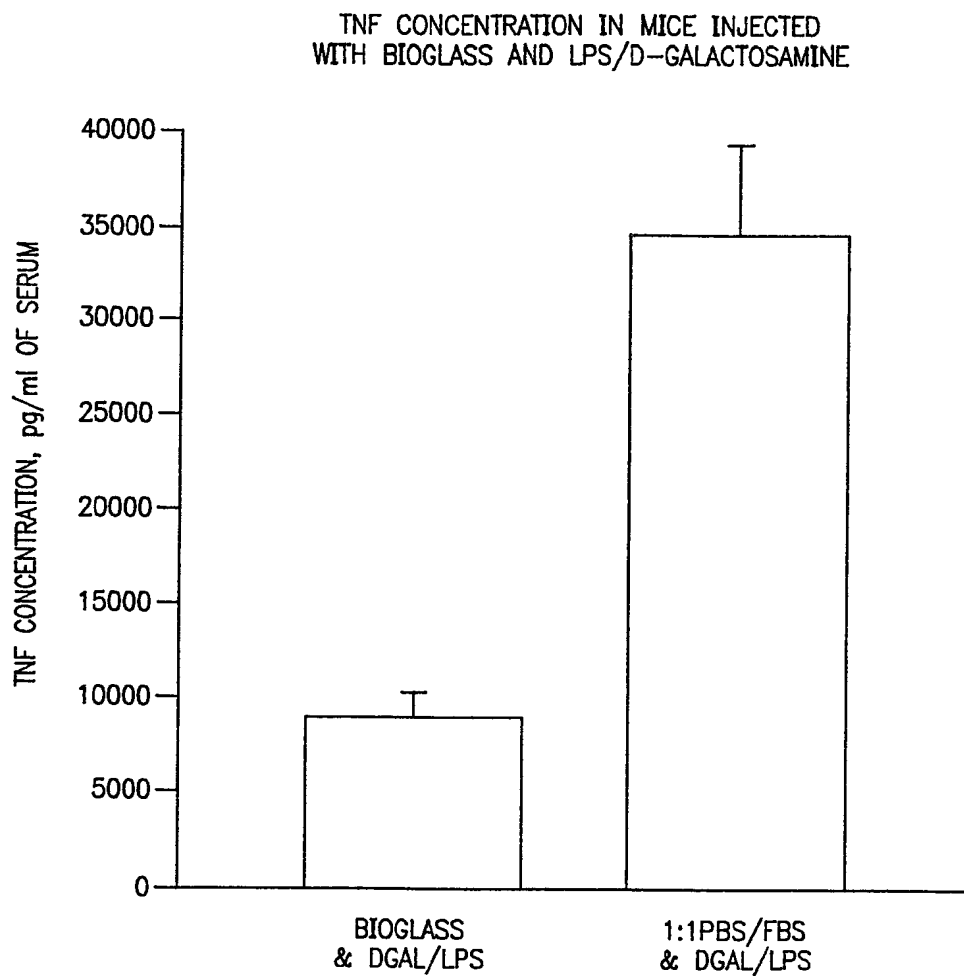


FIG. 5

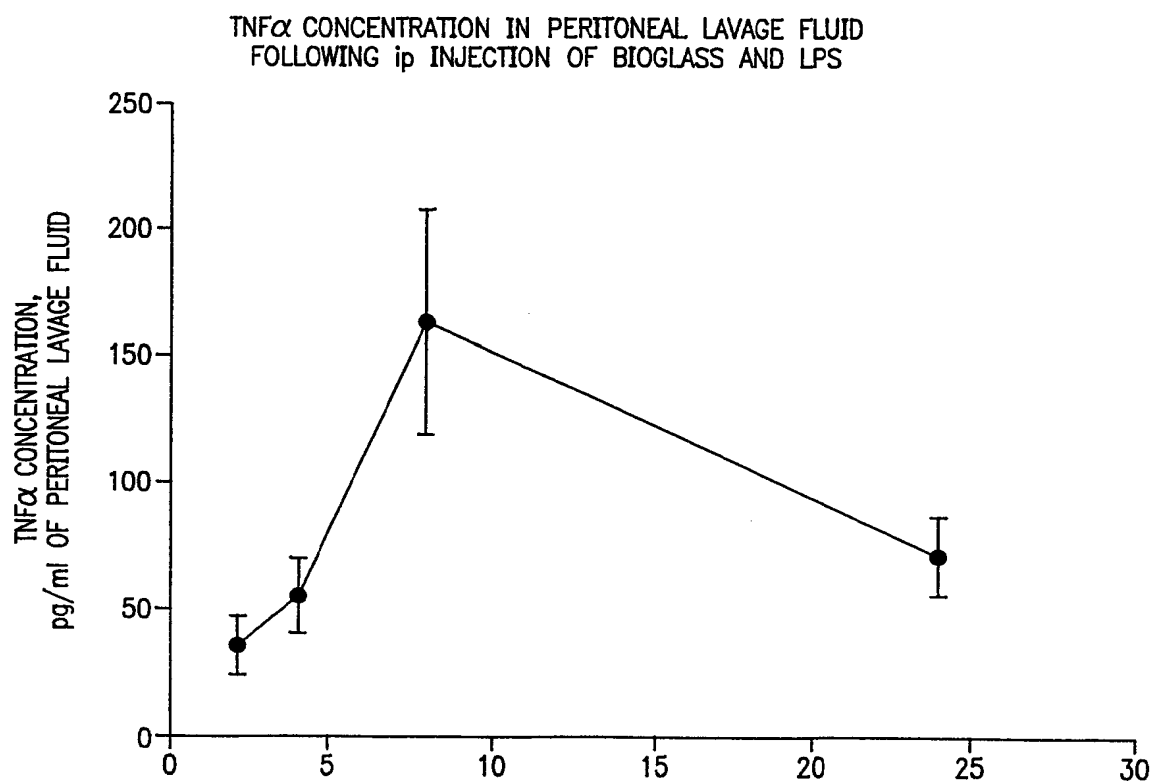


FIG. 6

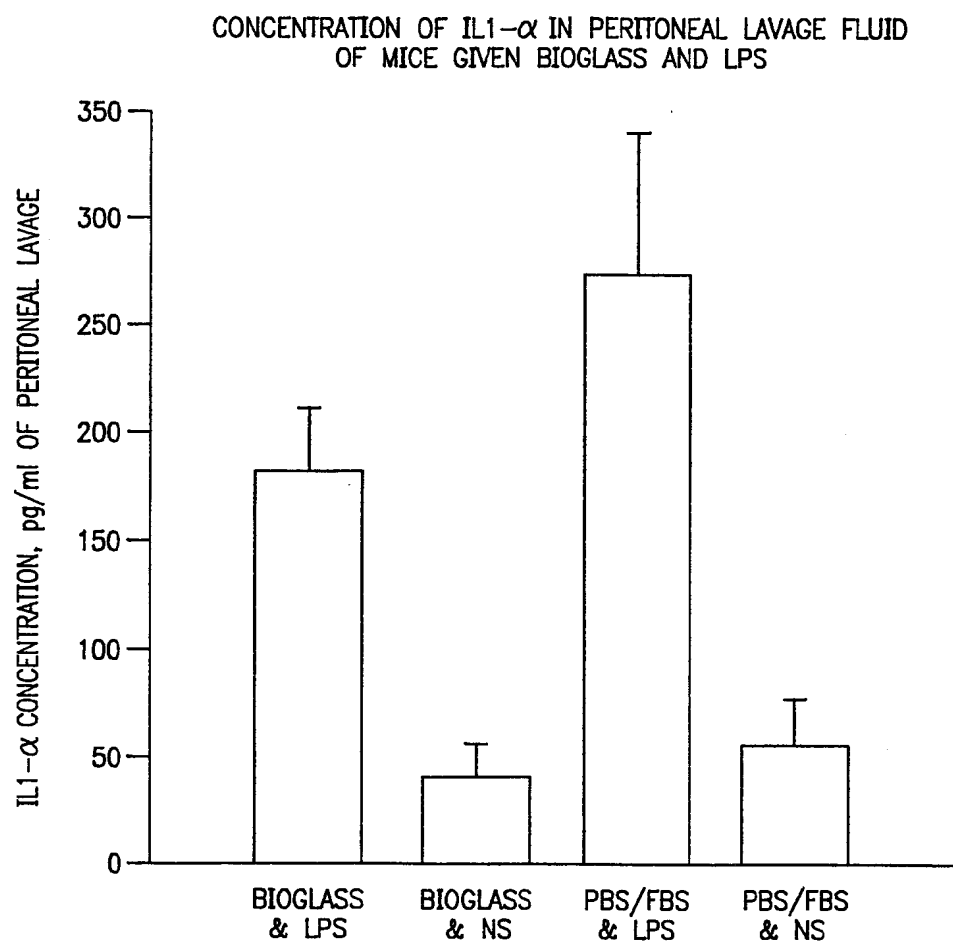


FIG. 7

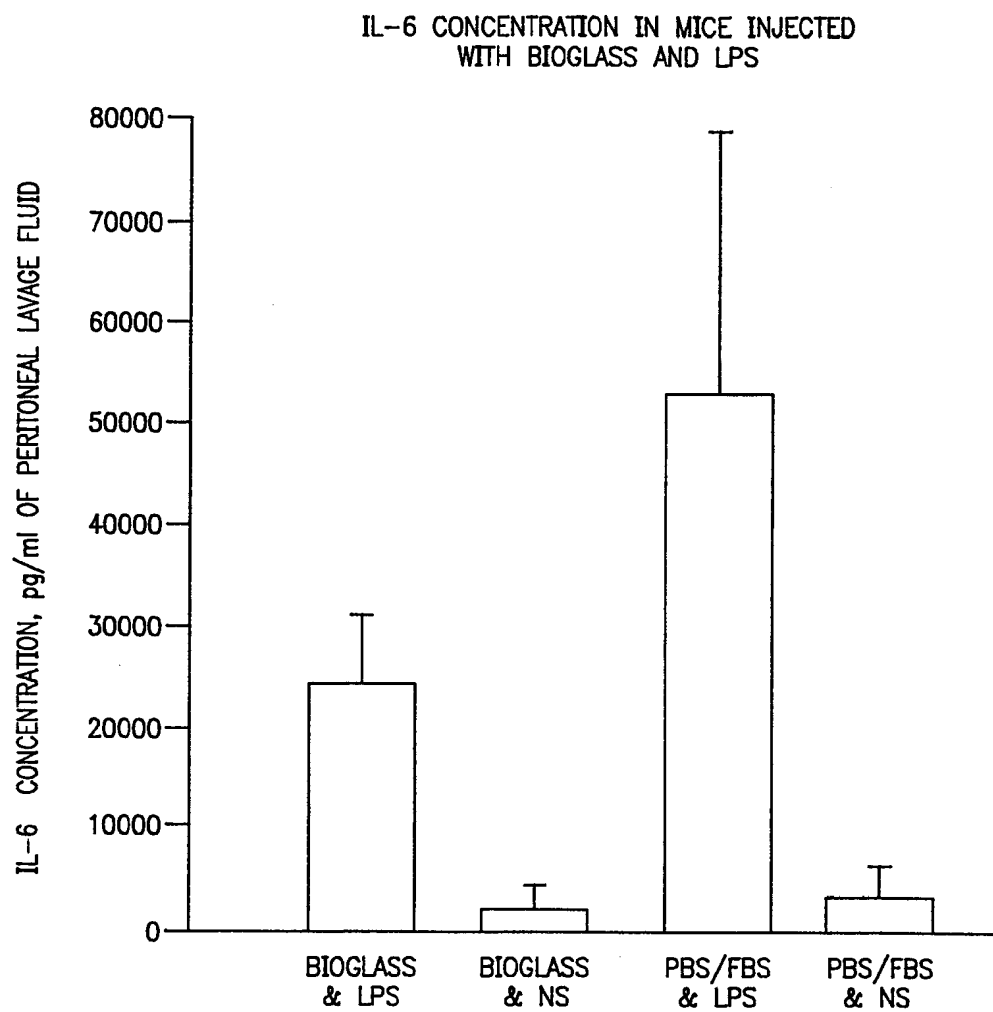


FIG. 8

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/11585

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 9/14

US CL : 424/489

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 424/489

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

MEDLINE, BIOSIS, CAPLUS, USPATFULL

search terms: glass particles, bioactive glass

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y,E	US 6,086,374 A (LITKOWSKI et al) 11 July 2000, see entire document.	1, 3-10
Y,P	US 5,990,380 A (MAROTTA et al) 23 November 1999, see entire document.	1-10
A,E	US 6,083,521 A (ACEMOGLU et al) 04 July 2000, see entire document.	11



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

25 AUGUST 2000

Date of mailing of the international search report

07 SEP 2000

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